

Document made available under the Patent Cooperation Treaty (PCT)

International application number: PCT/EP05/000556

International filing date: 18 January 2005 (18.01.2005)

Document type: Certified copy of priority document

Document details: Country/Office: EP
Number: 04447032.6
Filing date: 10 February 2004 (10.02.2004)

Date of receipt at the International Bureau: 25 April 2005 (25.04.2005)

Remark: Priority document submitted or transmitted to the International Bureau in compliance with Rule 17.1(a) or (b)



World Intellectual Property Organization (WIPO) - Geneva, Switzerland
Organisation Mondiale de la Propriété Intellectuelle (OMPI) - Genève, Suisse

18. 01. 2005

**Europäisches
Patentamt****European
Patent Office****Office européen
des brevets****Bescheinigung****Certificate****Attestation**

Die angehefteten Unterlagen stimmen mit der ursprünglich eingereichten Fassung der auf dem nächsten Blatt bezeichneten europäischen Patentanmeldung überein.

The attached documents are exact copies of the European patent application described on the following page, as originally filed.

Les documents fixés à cette attestation sont conformes à la version initialement déposée de la demande de brevet européen spécifiée à la page suivante.

Patentanmeldung Nr. Patent application No. Demande de brevet n°

04447032.6

Der Präsident des Europäischen Patentamts;
Im Auftrag

For the President of the European Patent Office

Le Président de l'Office européen des brevets
p.o.

R C van Dijk



Anmeldung Nr.:
Application no.: 04447032.6
Demande no:

Anmeldetag:
Date of filing: 10.02.04
Date de dépôt:

Anmelder/Applicant(s)/Demandeur(s):

Université Libre De Bruxelles
Avenue Franklin Roosevelt 50,
CP 161
1050 Brussels
BELGIQUE

Bezeichnung der Erfindung/Title of the invention/Titre de l'invention:
(Falls die Bezeichnung der Erfindung nicht angegeben ist, siehe Beschreibung.
If no title is shown please refer to the description.
Si aucun titre n'est indiqué se référer à la description.)

Phthalocyanine derivative layer in electronic multilayer devices and method for
the manufacturing thereof

In Anspruch genommene Priorität(en) / Priority(ies) claimed /Priorité(s)
revendiquée(s)
Staat/Tag/Aktenzeichen/State/Date/File no./Pays/Date/Numéro de dépôt:

Internationale Patentklassifikation/International Patent Classification/
Classification internationale des brevets:

C09K19/00

Am Anmeldetag benannte Vertragsstaaten/Contracting states designated at date of
filing/Etats contractants désignées lors du dépôt:

AT BE BG CH CY CZ DE DK EE ES FI FR GB GR HU IE IT LU MC NL
PT RO SE SI SK TR LI

PHTHALOCYANINE DERIVATIVE LAYER IN ELECTRONIC MULTILAYER
DEVICES AND METHOD FOR THE MANUFACTURING THEREOF

Field of the invention

[0001] The present invention is related to electronic devices and more particularly to a multilayer device comprising an organic active layer of a phthalocyanine derivative presenting a homeotropic alignment as well as a manufacturing method thereof.

State of the art

[0002] Liquid crystals (LC) are materials which can present a mesophase having long-range orientational order but no long-range positional order (Goodby J.W., Gray G.W., in Handbook of Liquid Crystals Demus D., Goodby J., Gray G.W., Spiess H.-W., Vill V. eds Wiley-VCH, 17-23). LC can be classified with respect to the shape of the molecules: calamitic LC have a rod or lath-like molecular structure while discotic LC have a relatively flat disc or sheet-shaped molecular structure.

[0003] The molecular structure of discotic LC confers them a unique combination of physical and chemical properties. The presence of the lateral chains grafted on the aromatic core induces a phase segregation, which leads to a large anisotropy of the material. In some cases, the discs stack on top of another to form columns, the different columns constituting a two-dimensional lattice. The latter can be hexagonal, rectangular, rectangular face

centered, oblique or tetragonal. The piling of the disc leads to the formation of a one-directional semi-conductive material. Indeed, in the columns, π - π stacking of aromatic cores allows electron hopping from one to another molecule, and the lateral chains play the role of insulator. Each column can thus be assimilated to an electric cable. Finally, discotic LC are very efficient in absorbing visible light.

[0004] Such properties make discotic liquid crystals very good candidates to build electronic devices like photovoltaic cells (PVCs), organic light emitting diodes (OLEDs), sensors, etc...

[0005] PVCs are designed to convert light into electricity. It requires at least two semi-conductive layers, one which is a negative charge carrier (n-type) and the other which is a positive charge carrier (p-type). Both layers can be constituted by LC, but the following pairs can also be considered: LC/semi-conducting polymer; LC/semi-conducting crystalline material, LC/semi-conducting amorphous material. Light adsorption leads to the formation of an excited state (exciton), which can only be efficiently split into a free charge pair at the interface between the two components. The charges can then migrate through the two layers to reach the electrodes, where they are collected.

[0006] In OLEDs, h^+ are injected into the p-type semi-conducting layer from a high work function electrode (indium tin oxide, gold, ...) and e^- are transferred into the p-type semi-conducting layer from a low work-function electrode (aluminum, silver, ...). The free charges have to be transferred into a third layer of electro-luminescent material, giving rise to charge recombination and light emission.

[0007] In liquid crystal-based sensors, a simple method to detect the molecule of interest is to measure a change in physical properties of the LC layer (texture, light absorption, mass) after the binding of the analyte to the mesogenic layer.

[0008] In all these applications, discotics have to be organized homeotropically, i.e. with the columns developing from one to the other layer between which they are sandwiched without any break (Fig. 1). From the electronic device point of view, homeotropic alignment increases exciton diffusion length up to 100 nm. In consequence, the probability that an exciton reach the interface between the two layers raise as well as the number of net charges created. From the sensor point of view, homeotropic alignment improves the sensitivity of the device.

[0009] In general, for nematic as well as for discotics, homeotropic alignment is induced by an alignment layer deposited on the electrode used in the electronic device. The said alignment layer can be obtained, for example, by an oblique evaporation process with SiO vapors (Lu et al, US patent 6,426,786) or by grafting of alkyl silane moieties with long alkyl chains (Kanbe et al, US patent 6,159,562). In some rare cases, spontaneous homeotropic alignment is observed between two substrate layers:

- 25 - a fluoroalkylated triphenylene derivative shows homeotropic alignment in the columnar hexagonal (Colhex) phase (around 130°C), between two indium tin oxide et soda-lime glass plates (Terasawa N. et al, ChemCom, 2003, 1678-1679);
- 30 - phthalocyanine lutetium and copper derivatives show homeotropic alignment in their columnar tetragonal (COLtet) phase (at 235° and 160°C, respectively) between two soda-lime glass and quartz substrates (Kazuaki H. et

al, Journal of Material Chemistry, 2001, 11, 423-433;
Kazuaki H. et al, Bull. Chem. Soc. Jpn, 2003, 76, 781-
787).

5 [0010] The present invention describes the spontaneous
homeotropic alignment of a phthalocyanine derivative
sandwiched between two substrate layers, at room
temperature and for a very wide range of substrate layers,
used as such.

10 Aims of the invention

[0011] The present invention aims to provide an
electronic device comprising a mesogenic active layer,
constituted by a phthalocyanine derivative, said
phthalocyanine derivative presenting a homeotropic
15 alignment at room temperature and for a very wide range of
adjacent substrate layers.

[0012] Furthermore, the present invention presents a
manufacturing method for electronic devices comprising a
mesogenic active layer of a phthalocyanine derivative.

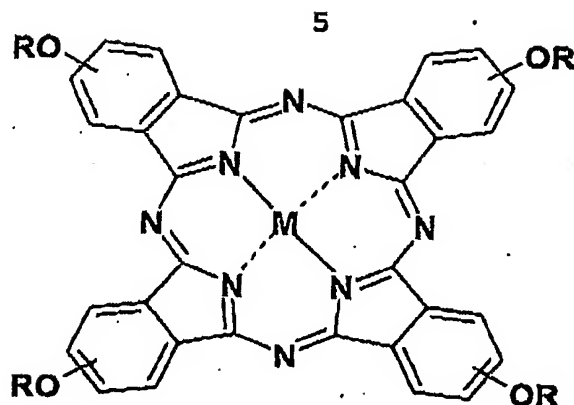
20

Summary of the invention

[0013] The present invention discloses an electronic
device including at least one organic semi-conducting layer
comprising a homeotropically organized phthalocyanine
25 derivative 1 sandwiched between at least two substrate
layers 2, 3.

[0014] In a first aspect of the present invention, the
phthalocyanine derivative is selected from the group
consisting of:

30



wherein: - M can be 2 atoms (Li_2 or H_2) or one metal atom (Zn, Cu, ...),

- R is an aliphatic moiety with the structure

$$\begin{array}{c} \text{CH}_2-\text{CH}-(\text{C}_n\text{H}_{n-2}) \\ | \\ (\text{C}_{n-2}\text{H}_{n-4}) \end{array}$$

with n going from 4 to 12

[0015] In a preferred aspect of the present invention, the substrate layers are inorganic layers.

[0016] Advantageously, the inorganic layers are soda lime glass, quartz or silicon.

[0017] In a general aspect of the present invention, the inorganic layers are coated with n-type liquid crystalline, crystalline or amorphous molecular semi-conductors selected from the group consisting of hexaazatriphenylene, hexaazatrinaphthalene, dodeca-azatri-phenylene, hexaazatri-isooxanaphthalene, hexa-azatri-isothianaphthalene, tricycloquinazoline, perylene derivatives and fullerene derivatives.

[0018] Advantageously, the substrate layers are organic polymer layer selected from the group consisting of polytetrafluoro-ethylène, polyethylene, polyethylene-terephthalate, polycarbonate, polyvinylchloride, polyurethane, poly-propylene, poly(methyl methacrylate).

[0019] Furthermore, the substrate layers further comprises a coating layer selected from the group consisting of metals or metal oxides.

[0020] In a particular aspect of the present invention, the metals or metal oxides are selected from the group consisting of silver, gold, aluminum, magnesium, indium tin oxide, tin oxide, zinc oxide, titanium oxide, gallium oxide, yttrium oxide and praseodymiumoxide.

[0021] In a particular aspect of the present invention, the substrate layers are coated with semi-conductive polymers selected from the group consisting of PEDOR, PEDOT-PSS, polyaniline, polypyrrole and polythiophène.

[0022] In an additional particular aspect of the present invention, the substrate layers or said coating layer are grafted by functional moiety's.

[0023] In another particular aspect of the present invention, the functional moiety's are silane groups.

[0024] Additionally, the present invention discloses a method for the manufacturing of the electronic device of the invention comprising the following steps:

- Depositing 1 to 5mg/cm² of a phthalocyanine derivative layer (1) on a first substrate layer (2) and covering said phthalocyanine derivative layer (1) fixed on the first substrate layer (2) by the second substrate layer (3) to build a sandwich.
- Heating the obtained sandwich in a hot plate at a temperature slightly above the isotropisation temperature,
- Applying a slight pressure on the second substrate layer (2) in order to make the film thinner,
- Cooling down the film at a cooling rate of 10°C/min to a temperature well below the isotropisation temperature in a stable position.

Short description of the drawings

[0025] Figure 1 is a schematic representation of homeotropic alignment of discotic molecules between two substrate layers.

[0026] Figure 2 is a schematic view of the electronic device with a phthalocyanine derivative layer 1 organized homeotropically and sandwiched in between a first 2 and a second substrate layer 3. Both layers being possibly coated with a coating layer 4.

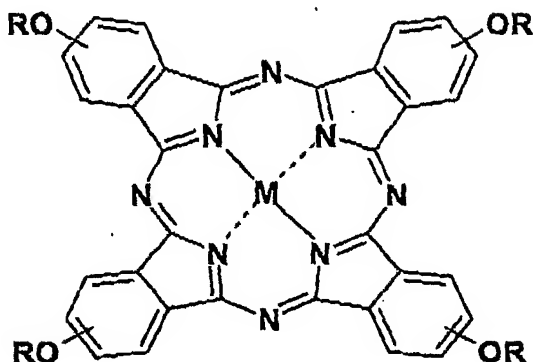
Detailed description of the invention

[0027] The present invention discloses an electronic device comprising, at least, one organic layer of a homeotropically organized phthalocyanine derivative sandwiched between a first and a second substrate layer 2 and 3.

[0028] The manufacturing method can be described by the following steps:

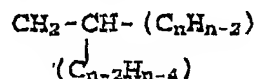
- 20 - Depositing 1 to .5mg/cm² of a phthalocyanine derivative layer 1 on a first substrate layer 2 and covering said phthalocyanine derivative layer 1 fixed on the first substrate layer 2 by the second substrate layer 3 to build a sandwich.
- 25 - Heating the obtained sandwich in a hot plate at a temperature slightly above the temperature at which the mesogenic become a liquid (isotropisation temperature).
- Applying a slight pressure on the second substrate layer 3 in order to obtain a film several microns thick.
- 30 Cooling down the film at a cooling rate of 10°C/min to a temperature well below the isotropisation temperature in a stable position in order not to loose the homeotropic alignment.

[0029] Phthalocyanine derivatives have the following chemical structure:



wherein: - M can be 2 atoms (Li_2 or H_2) or one metal atom (Zn, Cu, ...),

- R is an aliphatic moiety with the structure:



with n going from 4 to 12, the size of the chain thereof allowing the molecule to be liquid crystalline with an isotropic transition below the decomposition temperature and more specifically below 200°C . This temperature is the upper limit to be considered to keep reasonable conditions in the industrial process.

[0030] The substrate layers have to be stiff and flat surfaces like for example, soda lime glass, silicon, quartz,... In this case the substrate has to be carefully cleaned by a mixture of H_2O_2 and H_2SO_4 (1:10, v:v) before use. The low thickness of such substrate layers (100 μm) allowing a certain flexibility for a serie of applications.

[0031] Glass and related materials can advantageously be coated by metal or metal oxide in order to provide electrodes (2 and 3 coated by 4). Typical coating materials are the following: silver, gold, aluminium, magnesium,

indium tin oxide, tin oxide, zinc oxide, titanium oxide, gallium oxide, yttrium oxide praseodymium oxide.

[0032] In order to improve interactions between the substrate layers and the phthalocyanine derivative, both
5 glass (or related materials) substrates, coated or not with metal oxides can be grafted by alkylsilanes with short alkyl chains.

[0033] Glass and related materials can be replaced by polymer plates, coated or not with metal or metal oxide, in
10 order to allow for example, lamination of the devices on textiles. This operation consist to fix solar cell onto a textile under high pressure with a glue comprising a molten polymer. Without being limitative, good candidates for such substrates are the following: polytetrafluoroethylène,
15 polyethylene-(terephthalate), polycarbonate, polyvinyl-chloride, poly-urethane, polypropylene, poly(methyl methacrylate), ...

[0034] Glass and related materials can also be spin-coated (2 and 3 coated with 4) with n-type polymers, in
20 order to make the injection of charges in the system easier or in order to provide an electronic device like PVCs or OLEDs, which require the combination of materials with hole carriers properties (p-type, phthalocyanine derivative), one the one hand, and electron carriers properties on the
25 other hand. In this case, polymer candidates are the following:, PEDOT-PSS, polyoxadiazoles, poly(9,9-dioctylfluorene-co-benzothiadiazole), poly(9,9-dioctylfluorene), poly-pyridines, polyquinoxalines, polyquinolines, ...

30 [0035] Advantageously, semi-conductive polymer layer can be replaced by n-type liquid crystal, crystalline or amorphous molecular semi-conductor, deposited by spin-coating technique (2 or 3 coated with 4). Examples of such materials are: hexaazatriphenylene, hexaazatrinaphthalene,

dodecaazatriphenylene, hexaazatri-isooxanaphthalene, hexaazatriisothianaphthalene, tricyclo-guinazoline, perylene derivatives, fullerene derivatives and especially C61-butyric acid methyl ester.

5 Exemples

[0036] An electronic device comprising a phthalocyanine derivative layer with a lateral chain with $n = 12$, sandwiched between a first substrate layer of a glass substrate coated with Indium Tin Oxide (ITO) and a second substrate layer of a glass plate coated with aluminum and spin-coated with a C61-butyric acid methyl ester (PCBM) layer.

The device is obtained with the following manufacturing method:

- 15 - A glass plate 3 covered by an aluminum layer is spin-coated with a PCBM solution (4 mg/ml in toluene) at 1500 rpm with an acceleration of 1500 rpm/sec in order to deposit a thin PCBM layer 4 on the glass plate.
- 1-5 mg of a phthalocyanine derivative layer 1 is deposited on the ITO coated glass plate 2 and covered with the PCBM coated glass plate 3 + 4.
- 20 - The obtained sandwich is heated in a hot plate at 200°C, in order to obtain a liquid phthalocyanine derivative layer 1.
- 25 - A slight pressure is applied on the second substrate layer 3 + 4 in order to obtain a film of a phthalocyanine derivative layer 1 of several microns thick. The film is cooled down at a cooling rate of 10°C/min to ambient temperature in a stable position to avoid the loosing of
- 30 the homeotropic alignment.

[0037] Legend

1. Phthalocyanine derivative layer
2. First substrate layer

11

3. Second substrate layer

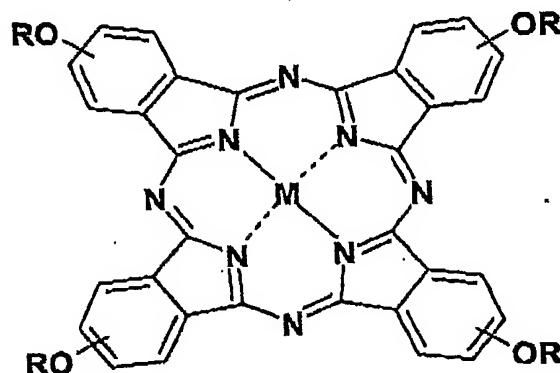
4. Coating layer

12

CLAIMS

1. Electronic device including at least one organic semi-conducting layer comprising a homeotropically organized phthalocyanine derivative (1) sandwiched between at least two substrate layers (2,3).

2. Electronic device according to Claim 1, wherein said phthalocyanine derivative is selected from the group consisting of:



wherein: - M can be 2 atoms (Li_2 or H_2) or one metal atom (Zn, Cu, ...),

- R is an aliphatic moiety with the structure

$$\begin{array}{c} \text{CH}_2-\text{CH}-(\text{C}_n\text{H}_{n-2}) \\ | \\ (\text{C}_{n-2}\text{H}_{n-4}) \end{array}$$

with n going from 4 to 12

3. Electronic device according to Claim 1, wherein said substrate layers are inorganic layers.

4. Electronic device according to Claim 3, wherein said inorganic layers are soda lime glass, quartz or silicon.

5. Electronic device according to Claim 3, wherein said inorganic layers are coated with n-type liquid crystalline, crystalline or amorphous molecular

+32 4 2229061

OFFICE VAN MALDEREN

10.FEV. 04 (MA) 14:02

+32 4 2229061

PAGE 20/24

semi-conductors selected from the group consisting of hexaazatriphenylene, hexaazatrinaphthalene, dodeca-azatriphenylene, hexaazatriisooxanaphthalene, hexa-azatriisothianaphthalene, tricycloquinazoline, perylene
5 derivatives and fullerene derivatives.

6. Electronic device according to Claim 1, wherein said substrate layers are organic polymer layer selected from the group consisting of polytetrafluoroethylene, polyethylene, polyethyleneterephthalate,
10 polycarbonate, polyvinylchloride, polyurethane, polypropylene, poly(methyl methacrylate).

7. Electronic device according to Claim 3 or 4, wherein said substrate layers further comprises a coating layer selected from the group consisting of metals
15 or metal oxides.

8. Electronic device according to Claim 5, wherein said metals or metal oxides are selected from the group consisting of silver, gold, aluminum, magnesium, indium tin oxide, tin oxide, zinc oxide, titanium oxide,
20 gallium oxide, yttrium oxide and praseodymiumoxide.

9. Electronic device according to Claim 3 or 4, wherein said substrate layers are coated with semi-conductive polymers selected from the group consisting of PEDOR, PEDOT-PSS, polyaniline, polypyrrole and
25 polythiophene.

10. Electronic device according to any of the preceding claims, wherein said substrate layers or said coating layer are grafted by functional moiety's.

11. Electronic device according to Claim
30 7, wherein said functional moiety's are silane groups.

12. Method for the manufacturing of the electronic device of Claim 1 comprising the following steps:

- Depositing 1 to 5mg/cm² of a phthalocyanine derivative layer (1) on a first substrate layer (2) and covering said phthalocyanine derivative layer (1) fixed on the first substrate layer (2) by the second substrate layer (3) to build a sandwich.
- Heating the obtained sandwich in a hot plate at a temperature slightly above the isotropisation temperature,
- Applying a slight pressure on the second substrate layer (2) in order to make the film thinner,
- Cooling down the film at a cooling rate of 10°C/min to a temperature well below the isotropisation temperature in a stable position.

13. Use of the electronic device of Claim 1 in photovoltaic cells, organic light emitting diodes and sensors.

ABSTRACTPHthalOCYANINE DERIVATIVE LAYERS IN ELECTRONIC MULTILAYER
DEVICES AND METHOD FOR THE MANUFACTURING THEREOF

5

The present invention relates to an electronic device including at least one organic semi-conducting layer comprising a homeotropically organized phthalocyanine derivative sandwiched between at least two substrate layers.

10

1/1

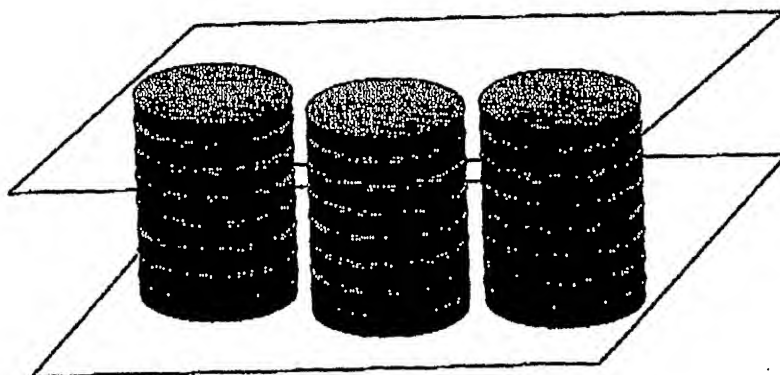


Fig. 1

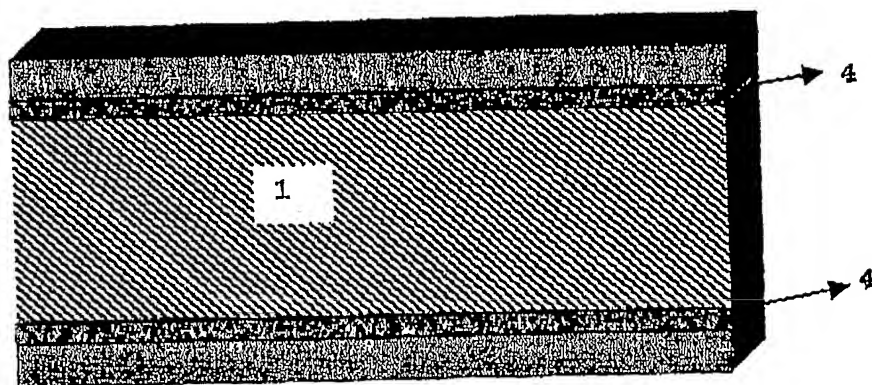


Fig. 2